

TITLE OF THE INVENTION

COMPOSITION FOR FORMING POROUS FILM, POROUS FILM AND
METHOD FOR FORMING THE SAME, INTERLEVEL INSULATOR FILM, AND
SEMICONDUCTOR DEVICE

5 CROSS-REFERENCE TO RELATED APPLICATION

The present application claims priority to Japanese Patent Application No. 2002-329124, filed November 13, 2002, the disclosure of which is incorporated herein by reference in its entirely.

10

BACKGROUND OF THE INVENTION

1. Field of the invention

The present invention relates to a composition for film formation, which can be formed into a porous film that excels in dielectric properties, adhesion, film consistency and mechanical strength, and has reduced absorption; a porous film and a method for forming the same; and a semiconductor device, which contains the porous film inside.

2. Description of the related art

20 In the fabrication of semiconductor integrated circuits, as the circuits are packed tighter, an increase in interconnection capacitance, which is a parasitic capacitance between metal interconnections, leads to an increase in interconnection delay time, thereby hindering the enhancement 25 of the performance of semiconductor circuits. The

interconnection delay time is called an RC delay, which is in proportion to the product of the electric resistance of the metal interconnections and the static capacitance between the interconnections. Reducing the interconnection delay time
5 requires reducing the resistance of metal interconnections or the interconnection capacitance.

The reduction in resistance of the interconnection metal and the interconnection capacitance can prevent a densely packed semiconductor device from causing an
10 interconnection delay, thereby realizing a smaller and faster semiconductor device with reduced power consumption.

In an attempt to reduce the resistance of metal interconnections, in recent years, metallic copper interconnections have been employed more than conventional
15 aluminum interconnections in the structure of a device. However, use of this structure all has limits in the enhancement of the performance, so the reduction in interconnection capacitance is an urgent necessity for higher performance of semiconductors.

20 One method for reducing interconnection capacitance is to reduce the relative permittivity (dielectric constant) of an interlevel insulator film disposed between metal interconnections. As such an insulator film with a low relative permittivity, it has been considered to use a porous
25 film instead of a silicon oxide film, which has been used

conventionally. A porous film can be said to be the only practical film as a material with a relative permittivity of 2.0 or less, and various methods for forming a porous film have been proposed

5 A first method for forming a porous film is as follows: a precursor solution of a siloxane polymer containing a thermally unstable organic component is synthesized; then the precursor solution is applied on the substrate to form a coating film; and later, a heat treatment is applied to
10 decompose and volatilize the organic component. The result is a number of micro-pores formed in the film.

As a second method for forming a porous film, it is well known to carry out processing as follows: a silica sol solution is applied onto a substrate by coating or using a
15 CVD method so as to form a wet gel; and then the silica sol is subjected to a condensation reaction while restricting volume reduction by controlling the speed of the evaporation of the solvent from the wet gel.

As a third method for forming a porous film, it is well
20 known that a silica micro-particle solution is applied on a substrate to form a coating film, and then the coating film is sintered to form a number of micro-pores between silica micro-particles.

As a fourth method, Japanese Patent Provisional
25 Publication No. 2000-44875 proposes a composition for porous

film formation, which is characterized by containing a compound having (A) a component expressed by $(R')_mSi(O R'')_{4-m}$ (R' and R'' are univalent organic radicals, and m is an integer of 0 to 2); (B) a metal chelate compound; and (C) a compound 5 having a polyalkylene oxide structure.

However, these methods have respective major drawbacks as follows.

In the first method for forming a porous film, the synthesis of the precursor solution of the siloxane polymer 10 increases the cost. In addition, the formation of the coating film by coating the precursor solution increases the amount of silanol groups remaining in the coating film, which causes a degassing phenomenon indicating the evaporation of water and the like in the heat treatment process that is 15 conducted later and, which also deteriorates the film quality due to the porous film absorbing humidity.

In the second method for forming a porous film, the speed control of the evaporation of the solvent from the wet gel requires a special type of coating device, which 20 increases the cost. In addition, a significant amount of silanol remains on the surface of the micro-pores, which must be silanized because otherwise hygroscopicity is high and the film quality decreases. The silanization makes the process more complicated. In the case where a wet gel is formed by 25 the CVD process, it is necessary to use a special type of CVD

device, which is different from the plasma CVD device generally used in the semiconductor process, thereby also increasing the cost.

In the third method for forming a porous film, the
5 diameter of the micro-pores formed between the silica micro-
particles, which is determined by the accumulation structure
of the silica micro-particles that are accumulated
geometrically, becomes very large. This makes it difficult to
set the relative permittivity of the porous film to 2 or
10 below.

In the case of the fourth method, out of the three components (A), (B), and (C), the metal chelate compound of (B) is essential to increase the compatibility of the components (A) and (C), and to make the thickness of the
15 coating film uniform after being hardened. However, it is not preferable because it makes the manufacturing process complicated and increases the cost. Therefore, it is desired to develop a material, which enables a homogeneous solution to be formed without a chelate component and the coating film
20 to be flat after being hardened.

Contrary to the conventional method for forming porous film, it has been found that aluminosilicate, silica and the like are first condensed using a mold of micelle formed by a surfactant and then the surfactant is removed from the
25 produced construction by calcination or solvent extraction,

which results in the formation of a porous structure having a mesopore (2-50 nm in diameter)-sized channel structure. For instance, according to Inagaki et al. (J. Chem. Soc. Chem. Commun., p. 680, 1993), polysilicate is reacted water in the presence of surfactant as a mold. Moreover, according to the Japanese Patent Provisional Publication No. 9-194298, the tetraalkoxysilane is reacted in water under an acidic condition in the presence of surfactant as a mold and coated on a substrate to form a silica porous film containing pores having pore diameter of 1 to 2 nm.

However, there is a problem with these methods. In the former, although a powdery porous body is easily made, a porous film cannot be formed on a substrate which is used for producing a semiconductor device. In the latter, although a porous film can be formed, the orientation of pores cannot be controlled so that uniform film in a large area thereof cannot be obtained.

Further, according to Japanese Patent Provisional Publication No. 2001-130911, the silica meso-porous film is formed using a mixture of a surfactant and a product obtained by acid hydrolysis and condensation of silicon alkoxide, where the mixture has been prepared and stabilized at pH value of 3 or less.

However, in this case, because a solute concentration is specified, it is difficult to control the coating film

thickness. Then, it is difficult to apply this method to an actual semiconductor manufacturing process. Moreover, when this solution is diluted with water, although the thickness of the coated film can be controlled, the rate of
5 condensation polymerization of the silica component increases, so that the stability of the coating liquid is lost.

On the other hand, the following methods are in the 2001-110529th according to the Japanese Patent Provisional Publication Nos. 2001-110529 and 2001-203197, a coating
10 liquid is obtained by hydrolysis and condensation of a silane compound and provides excellent dielectric properties. Because a modulus of elasticity required in the manufacturing process of an actual semiconductor is 5 GPa or higher, however, the mechanical strength is not sufficient in these
15 inventions.

As mentioned above, when porous film produced by conventional material is used as an insulator film in multi-level interconnects of the semiconductor device, there are problems such that moisture-absorbed porous film increases dielectric
20 constant and lowers the adhesiveness in the lamination of films. Moreover, when the dielectric constant of the porous film used as an insulator film increases, the RC delay in the multi-level interconnects of the semiconductor device is increased. Consequently, the performance of the
25 semiconductor device (high speed and low power consumption)

has not been improved. Furthermore, higher cost for forming porous film leads to higher cost for a semiconductor device. Moreover, when conventional porous film is incorporated as an insulator film into the multi-level interconnects of the
5 semiconductor device, there is a problem of the film having insufficient mechanical strength. The low mechanical strength results from the porous film property. Thus, there is a problem that low mechanical strength of the porous film as an insulator film lowers the reliability of a
10 semiconductor device.

SUMMARY OF THE INVENTION

An object of the invention is to provide a composition for forming film which can form a porous film excelling in dielectric constant, adhesiveness, uniformity of the film,
15 mechanical strength and having low hygroscopicity. Moreover, an object of the invention is to provide a high-performing and highly reliable semiconductor device comprising the porous film inside.

20 The inventors focused intently on the development of a coating liquid for forming the porous film. As a result, they reached the composition for forming porous film having low dielectric constant as well as high mechanical strength which is applicable to the semiconductor production process,
25 and the method for forming the porous film, based on that

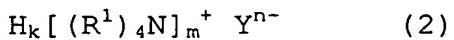
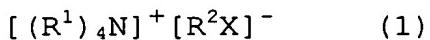
siloxane polymer is heated with so-called a structure-directing agent. Then, they completed the invention.

For example, according to the description of page 95 of "Creation and application of the molecular coordination material", supervised by Kunihiro Ichimura, published by CMC Publishing Co., Ltd., structure-directing agents generally refer to organic cation which is used in synthesis of zeolite and are known to play an important role in hydrothermal synthesis of high silica zeolite. A quaternary ammonium compound, especially hydroxide, is often preferably used as organic cation substance. Particularly in the field of manufacturing an insulator film having a low dielectric constant to which the invention relates, according to the Japanese Patent Provisional Publication No. 2002-30249, a method for manufacturing a insulator film having low dielectric constant using quaternary ammonium hydroxide or quaternary ammonium halide is known.

Because the quaternary ammonium hydroxide is alkaline, however, the stability of the produced coating solution decreases markedly, making practical use very difficult. Moreover, when quaternary ammonium halide is used, the residual halogen in the insular film which has been coated and sintered makes lead wires in circuits corroded, thereby obstructing continuity and developing into the possible deterioration of semiconductor devices. Thus, these

substances are not suitable.

Therefore, according to the invention, provided is a composition for forming porous film, the composition comprising siloxane polymer and one or more quaternary ammonium salts represented by following formula (1) or (2):



wherein R^1 independently represents a straight chain or branched alkyl or aryl group having 1 to 10 carbons which may have a substituent and R^1 's may be same or different; R^2 represents a hydrogen atom or an straight chain or branched alkyl or aryl group having 1 to 10 carbons which may have a substituent; X represents CO_2 , OSO_3 or SO_3 ; Y represents SO_4 , SO_3 , CO_3 , O_2C-CO_2 , NO_3 or NO_2 ; and k is 0 or 1, m is 1 or 2 and n is 1 or 2 in proviso that $n=1$ requires $k=0$ and $m=1$, and $n=2$ requires $k=0$ and $m=2$, or $k=1$ and $m=1$.

The structure-directing agents conventionally known are quaternary ammonium hydroxide and quaternary ammonium halide. The quaternary ammonium hydroxide is a nucleation crystal growth model as indicated in P.P.E,A.D. Moor et al., Chem. Eur. J., 5, 7, 2083 (1998) so that the quaternary ammonium hydrolyzes and condensates silane monomers to form an aggregate from which zeolite crystal grows. Therefore, the quaternary ammonium hydroxide has ability to direct a structure as monomers grow to be crystal.

On the other hand, the structure-directing agent used for this invention has the effect of structure-directing by means of not using monomers but changing the configuration of polymer whose frame has already been formed. Actually, it is 5 supported by the GPC measurements of no increase in molecular weight distribution before and after addition of this structure-directing agent.

Such a quaternary ammonium salt compound is not known to have a property of a structure-directing agent until we 10 discovers. We has found the novel property and completed the invention.

Moreover, quaternary ammonium halide is conventionally known as a substance to form a zeolite structure or honeycomb structure by forming micelle between polymer and a quaternary 15 ammonium salt, in a similar manner to that of the invention.

On the other hand, the quaternary ammonium salt used for the invention is easily decomposed by heating for hardening the coated film so that problems such as corrosion of circuit do not occur. This is also different from conventional halide.

20 According to the invention, a method for forming a porous film comprising a step of applying the composition on a substrate, subsequently a drying step and a heating step for hardening the porous film. Moreover, a porous film obtainable from the composition for forming a porous film is 25 provided. These are applicable to the semiconductor

manufacturing process and provide a porous film having an excellent dielectric constant and mechanical properties.

According to the invention, the semiconductor device containing an internal porous film formable by the 5 composition for forming a porous film is provided. More specifically, a porous film is used as an insulator film in the multi-level interconnects in a semiconductor device. Consequently, because the mechanical strength of the semiconductor device is secured and the hygroscopic property 10 of the porous film is decreased, a semiconductor device containing an internal porous film having low dielectric constant is obtained. Because of lower dielectric constant of the insulator film, the parasitic capacitance of the area around the multi-level interconnects is decreased, leading to 15 the high-speed operation and low power consumption of the semiconductor device. Moreover, it is preferable in the semiconductor device of the invention that said porous film is between metal interconnections in a same layer of multi-level interconnects, or is between upper and lower metal 20 interconnection layers. This arrangement can achieve a high-performing and highly reliable semiconductor device.

The use of the composition for forming porous film of the invention can facilitate the formation of porous film having a desirably controlled thickness. This porous film 25 has low dielectric constant and excels in adhesion, film

uniformity and mechanical strength. In addition, use of the porous film formable by the composition of the invention as the insulator film of the multi-level interconnects can achieve a high-performing and highly reliable semiconductor
5 device.

BRIEF DESCRIPTION OF THE DRAWING

Fig. 1 is a schematic cross-sectional view of a semiconductor device of the invention.

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DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The siloxane polymer used for the invention can be a polymer obtained by a well-known method. For instance, the
15 siloxane polymer may be a polymer obtainable by the method introduced by C.J. Brinker and G.W. Scherer, "Sol-gel Science," Academic Press (1990); Kunio Ito, "Silicone Handbook," Nikkan Kogyo Shimbun Co. (1990); Sumio Sakka, "Science of the Sol-gel Method," Agne Shofu-sha (1988); or
20 Sumio Sakka, "Application of the Sol-gel Method" Agne Shofu-sha (1997).

Specifically, the siloxane polymer may be produced by hydrolysis and condensation of one or more silane compounds in the presence of acidic or basic catalyst in an organic
25 solvent to which water has been added, wherein the silane

compound comprises one or more hydrolysable substituents within the molecule.

The silane compound may include, but is not limited to, tetramethoxysilane, tetraethoxysilane, tetrapropoxysilane, 5 tetrabutoxysilane, methyltrimethoxysilane, methyltriethoxysilane, methyltripropoxysilane, ethyltrimethoxysilane, propyltrimethoxysilane, butyltrimethoxysilane, pentyltrimethoxysilane, hexyltrimethoxysilane, 2-ethylhexyltrimethoxysilane, 10 phenyltrimethoxysilane, dimethyldimethoxysilane, dimethyldiethoxysilane, trimethylmethoxysilane, triethylmethoxysilane, butyldimethylmethoxysilane, hexamethoxydisilane, hexaethoxydisilane, 1,2-bis(trimethoxysilyl)ethane, 1,2-bis(triethoxysilyl)ethane, 15 1,6-bis(trimethoxysilyl)hexane, 1,6-bis(triethoxysilyl)hexane, 1,4-bis(trimethoxysilyl)benzene, 1,4-bis(triethoxysilyl)benzene, 1,2-bis(dimethoxymethylsilyl)ethane, 1,2-bis(diethoxymethylsilyl)ethane, 1,6-bis(dimethoxymethylsilyl)hexane, 20 1,6-bis(diethoxymethylsilyl)hexane, 1,4-bis(dimethoxymethylsilyl)benzene, 1,4-bis(diethoxymethylsilyl)benzene, 1,2-bis(methoxydimethylsilyl)ethane, 1,2-bis(ethoxydimethylsilyl)ethane, 1,6-bis(methoxydimethylsilyl)hexane, 25 1,6-

bis(ethoxydimethylsilyl)hexane, 1,4-
bis(methoxydimethylsilyl)benzene and 1,4-
bis(ethoxydimethylsilyl)benzene.

The organic solvent used in the invention may include
5 methanol, ethanol, isopropyl alcohol, butanol, propylene
glycol monomethyl ether, propylene glycol monopropyl ether,
propylene glycol monopropyl ether acetate, ethyl lactate and
cyclohexanone. The solvent other than water may be added in
an amount of preferably 0.1 to 500 times, more preferably 1
10 to 100 times the weight of the silicon compound. The water
for the hydrolysis may be added in an amount of preferably
0.5 to 10 times, more preferably 1.0 to 4.0 times the mole
number required for complete hydrolysis of the silane
compound.

15 An example of the acid catalyst used at this time may
include inorganic acid such as hydrochloric acid, sulfuric
acid and nitric acid; sulfonic acid such as methanesulfonic
acid, benzenesulfonic acid, p-toluenesulfonic acid and
trifluoromethanesulfonic acid; organic acid such as formic
20 acid, acetic acid, propionic acid, oxalic acid, malonic acid,
fumaric acid, malenic acid, tartaric acid, citric acid and
malic acid; and phosphoric acid. The acid catalyst may be
added preferably 0.001 to 10 times, more preferably 0.01 to
1.0 times the mole number of the starting silicon compound.

25 On the other hand, basic catalyst which can be used in

this case may include ammonia; amine such as ethylamine, propylamine, diisopropylamine, triethylamine and triethanolamine; and alkali metal hydroxide or alkaline earth metal hydroxide such as sodium hydroxide, potassium hydroxide, 5 and calcium hydroxide. The basic catalyst may be added in an amount of preferably 0.001 to 10 times, more preferably 0.01 to 1.0 times the mole number of the starting silane compound.

The typical production procedure of a siloxane polymer may be as follows. The abovementioned organic solvent may be 10 mixed with water and the reaction catalyst. The silicon compound in a required amount may be then added thereto and treated at the prescribed temperature for a given time to produce target polymer. Physical properties of the obtained polymer can be changed depending on reaction temperature, an 15 amount of water to be added, a type of organic solvent used for the reaction, and concentration and type of the starting silicon compound.

The reaction temperature may be typically from 0°C to the boiling point of alcohol which is produced as the result 20 of the hydrolysis and condensation. It may be preferably from room temperature to 100°C. The reaction time may not be particularly limited. It may be typically between 10 minutes and 30 hours, more preferably 30 minutes to 10 hours.

The preferably weight-average molecular weight of the 25 siloxane polymer may be 10,000 to 1,000,000 using

polyethylene as standard with Gel Permeation Chromatograph (GPC). It may be because the property of porous film useful also as an insulation film a semiconductor device is realized.

To the obtained solution of siloxane polymer is added
5 quaternary ammonium salt represented by formula (1) or (2)
and treated. Accordingly, the polymer frame can be
transformed into a structure similar to the zeolite structure.

The R¹ represented by formulas (1) and (2) may include,
but not limited to, methyl, ethyl, n-propyl, iso-propyl, n-
10 butyl isobutyl, sec-butyl, n-pentyl, 2-methylbutyl, 3-
methylbuty, 2,2-dimethylpropyl, n-hexyl, 2-methylpentyl, 3-
methylpentyl, 4-methylpenty, 2,2-dimethylbutyl, 3,3-
dimethylbutyl, n-heptyl, 2-methylhexyl, 3-methylhexyl, 4-
methylhexyl, 5-methylhexyl, 2,2-dimethylpentyl, 3,3-
15 dimethylpentyl, 4,4-dimethylpentyl, 3-ethylpentyl, 4-
ethylpentyl, n-octyl, 2-methylheptyl, 3-methylheptyl, 4-
methylheptyl, 5-methylheptyl, 6-methylheptyl, 2,2-
dimethylhexyl, 3,3-dimethylhexyl, 4,4-dimethylhexyl, 5,5-
dimethylhexyl, 3-ethylhexyl, 4-ethylhexyl, 5-ethylhexyl, n-
20 nonyl, 2-methyloctyl, 3-methyloctyl, 4-methyloctyl, 5-
methyloctyl, 6-methyloctyl, 7-methyloctyl, 2,2-dimethylheptyl,
3,3-dimethylheptyl, 4,4-dimethylheptyl, 5,5-dimethylheptyl, 6,
6-dimethylheptyl, 3-ethylheptyl, 4-ethylheptyl, 5-ethylheptyl,
6-ethylheptyl, n-decyl, 2-methylnonyl, 3-methylnonyl, 4-
25 methylnonyl, 5-methylnonyl, 6-methylnonyl, 7-methylnonyl,

2,2-dimethyloctyl, 3,3-dimethyloctyl, 4,4-dimethyloctyl, 5,5-dimethyloctyl, 6,6-dimethyloctyl, 7,7-dimethyloctyl, 3-ethyloctyl, 4-ethyloctyl, 5-ethyloctyl, 6-ethyloctyl, 7-ethyloctyl, phenyl, o-tolyl, m-tolyl and p-tolyl.

5 The R² represented by formula (1) may include a hydrogen atom, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, n-pentyl, 2-methylbutyl, 3-methylbutyl, 2,2-dimethylpropyl, n-hexyl, 2-methylpentyl, 3-methylpentyl, 4-methylpentyl, 2,2-dimethylbutyl, 3,3-dimethylbutyl, n-heptyl, 10 2-methylhexyl, 3-methylhexyl, 4-methylhexyl, 5-methylhexyl, 2,2-dimethylpentyl, 3,3-dimethylpentyl, 4,4-dimethylpentyl, 3-ethylpentyl, 4-ethylpentyl, n-octyl, 2-methylheptyl, 3-methylheptyl, 4-methylheptyl, 5-methylheptyl, 6-methylheptyl, 2,2-dimethylhexyl, 3,3-dimethylhexyl, 4,4-dimethylhexyl, 5,5-dimethylhexyl, 3-ethylhexyl, 4-ethylhexyl, 5-ethylhexyl, n-nonyl, 2-methyloctyl, 3-methyloctyl, 4-methyloctyl, 5-methyloctyl, 6-methyloctyl, 7-methyloctyl, 2,2-dimethylheptyl, 15 3,3-dimethylheptyl, 4,4-dimethylheptyl, 5,5-dimethylheptyl, 6,6-dimethylheptyl, 3-ethylheptyl, 4-ethylheptyl, 5-ethylheptyl, 6-ethylheptyl, n-decyl, 2-methylnonyl, 3-methylnonyl, 4-methylnonyl, 5-methylnonyl, 6-methylnonyl, 7-methylnonyl, 2,2-dimethyloctyl, 3,3-dimethyloctyl, 4,4-dimethyloctyl, 5,5-dimethyloctyl, 6,6-dimethyloctyl, 7,7-dimethyloctyl, 3-ethyloctyl, 4-ethyloctyl, 5-ethyloctyl, 6-ethyloctyl, 18 7-ethyloctyl, phenyl, o-tolyl, m-tolyl and p-

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tolyl. In addition, one or more hydrogen atoms of the above example may be substituted by hydroxy, methoxy, ethoxy, n-propoxy, isopropoxy, phenoxy, methylthio, ethylthio, amino, methylamino dimethylamino, carboxy, methoxycarbonyl,
5 ethoxycarbonyl, formyloxy, acetoxy, benzoyloxy, formyl, acetyl or propionyl.

The X in formulation (1) represents CO₂, OSO₃ or SO₃.

The quaternary ammonium salt represented by formula (1) may include tetramethylammonium formate, tetramethylammonium acetate, tetramethylammonium propionate, tetramethylammonium butyrate, tetramethylammonium isobutyrate, tetramethylammonium valerate, tetramethylammonium pivalate, tetramethylammonium hexanoate, tetramethylammonium octanoate, tetramethylammonium decanoate, tetramethylammonium lactate,
15 trimethylethylammonium formate, trimethylethylammonium acetate, trimethylethylammonium propionate, trimethylethylammonium butyrate, trimethylethylammonium isobutyrate, trimethylethylammonium valerate, trimethylethylammonium pivalate, trimethylethylammonium hexanoate, trimethylethylammonium octanoate,
20 trimethylethylammonium decanoate, trimethylethylammonium lactate, trimethylpropylammonium formate, trimethylpropylammonium acetate, trimethylpropylammonium propionate, trimethylpropylammonium butyrate,
25 trimethylpropylammonium isobutyrate, trimethylpropylammonium

valerate, trimethylpropylammonium pivalate,
trimethylpropylammonium hexanoate, trimethylpropylammonium
octanoate, trimethylpropylammonium decanoate,
trimethylpropylammonium lactate, trimethylbutylammonium
5 formate, trimethylbutylammonium acetate,
trimethylbutylammonium propionate, trimethylbutylammonium
butyrate, trimethylbutylammonium isobutyrate,
trimethylbutylammonium valerate, trimethylbutylammonium
pivalate, trimethylbutylammonium hexanoate,
10 trimethylbutylammonium octanoate, trimethylbutylammonium
decanoate, trimethylbutylammonium lactate,
trimethylpentylammonium formate, trimethylpentylammonium
acetate, trimethylpentylammonium propionate,
trimethylpentylammonium butyrate, trimethylpentylammonium
15 isobutyrate, trimethylpentylammonium valerate,
trimethylpentylammonium pivalate, trimethylpentylammonium
hexanoate, trimethylpentylammonium octanoate,
trimethylpentylammonium decanoate, trimethylpentylammonium
lactate, trimethylhexylammonium formate,
20 trimethylhexylammonium acetate, trimethylhexylammonium
propionate, trimethylhexylammonium butyrate,
trimethylhexylammonium isobutyrate, trimethylhexylammonium
valerate, trimethylhexylammonium pivalate,
trimethylhexylammonium hexanoate, trimethylhexylammonium
25 octanoate, trimethylhexylammonium decanoate,

trimethylhexylammonium lactate, trimethylheptylammonium
formate, trimethylheptylammonium acetate,
trimethylheptylammonium propionate, trimethylheptylammonium
butyrate, trimethylheptylammonium isobutyrate,
5 trimethylheptylammonium valerate, trimethylheptylammonium
pivalate, trimethylheptylammonium hexanoate,
trimethylheptylammonium octanoate, trimethylheptylammonium
decanoate, trimethylheptylammonium lactate,
trimethyloctylammonium formate, trimethyloctylammonium
10 acetate, trimethyloctylammonium propionate,
trimethyloctylammonium butyrate, trimethyloctylammonium
isobutyrate, trimethyloctylammonium valerate,
trimethyloctylammonium pivalate, trimethyloctylammonium
hexanoate, trimethyloctylammonium octanoate,
15 trimethyloctylammonium decanoate, trimethyloctylammonium
lactate, trimethylnonylammonium formate,
trimethylnonylammonium acetate, trimethylnonylammonium
propionate, trimethylnonylammonium butyrate,
trimethylnonylammonium isobutyrate, trimethylnonylammonium
20 valerate, trimethylnonylammonium pivalate,
trimethylnonylammonium hexanoate, trimethylnonylammonium
octanoate, trimethylnonylammonium decanoate,
trimethylnonylammonium lactate, trimethyldecylammonium
formate, trimethyldecylammonium acetate,
25 trimethyldecylammonium propionate, trimethyldecylammonium

butyrate, trimethyldecylammonium isobutyrate,
trimethyldecylammonium valerate, trimethyldecylammonium
pivalate, trimethyldecylammonium hexanoate,
trimethyldecylammonium octanoate, trimethyldecylammonium
5 decanoate, trimethyldecylammonium lactate,
methyltriethylammonium formate, methyltriethylammonium
acetate, methyltriethylammonium propionate,
methyltriethylammonium butyrate, methyltriethylammonium
isobutyrate, methyltriethylammonium valerate,
10 methyltriethylammonium pivalate, methyltriethylammonium
hexanoate, methyltriethylammonium octanoate,
methyltriethylammonium decanoate, methyltriethylammonium
lactate, methyltripropylammonium formate,
methyltripropylammonium acetate, methyltripropylammonium
15 propionate, methyltripropylammonium butyrate,
methyltripropylammonium isobutyrate, methyltripropylammonium
valerate, tripropylammonium pivalate, methyltripropylammonium
hexanoate, methyltripropylammonium octanoate,
methyltripropylammonium decanoate, methyltripropylammonium
20 lactate, methyltributylammonium formate,
methyltributylammonium acetate, methyltributylammonium
propionate, methyltributylammonium butyrate,
methyltributylammonium isobutyrate, methyltributylammonium
valerate, methyltributylammonium pivalate,
25 methyltributylammonium hexanoate, methyltributylammonium

octanoate, methyltributylammonium decanoate,
methyltributylammonium lactate, methyltripentylammonium
formate, methyltripentylammonium acetate,
methyltripentylammonium propionate, methyltripentylammonium
5 butyrate, methyltripentylammonium isobutyrate,
methyltripentylammonium valerate, methyltripentylammonium
pivalate, methyltripentylammonium hexanoate,
methyltripentylammonium octanoate, methyltripentylammonium
decanoate, methyltripentylammonium lactate, methyltrihexyl
10 ammonium formate, methyltrihexylammonium acetate,
methyltrihexylammonium propionate, methyltrihexylammonium
butyrate, methyltrihexylammonium isobutyrate,
methyltrihexylammonium valerate, methyltrihexylammonium
pivalate, methyltrihexylammonium hexanoate,
15 methyltrihexylammonium octanoate, methyltrihexylammonium
decanoate, methyltrihexylammonium lactate,
methyltriheptylammonium formate, methyltriheptylammonium
acetate, methyltriheptylammonium proionate,
methyltriheptylammonium butyrate, methyltriheptylammonium
20 isobutyrate, methyltriheptylammonium valerate,
methyltriheptylammonium pivalate, methyltriheptylammonium
hexanoate, methyltriheptylammonium octanoate,
methyltriheptylammonium decanoate, methyltriheptylammonium
lactate, methyltrioctylammonium foramtne,
25 methyltrioctylammonium acetate, methyltrioctylammonium

propionate, methyltrioctylammonium butyrate,
methyltrioctylammonium isobutyrate, methyltrioctylammonium
valerate, methyltrioctylammonium pivalate,
methyltrioctylammonium hexanoate, methyltrioctylammonium
5 octanoate, methyltrioctylammonium decanoate,
methyltrioctylammonium lactate, methyltrinonylammonium
formate, methyltrinonylammonium acetate,
methyltrinonylammonium propionate, methyltrinonylammonium
butyrate, methyltrinonylammonium isobutyrate,
10 methyltrinonylammonium valerate, methyltrinonylammonium
pivalate, methyltrinonylammonium hexanoate,
methyltrinonylammonium octanoate, methyltrinonylammonium
decanoate, methyltrinonylammonium lactate,
methyltridecylammonium formate, methyltridecylammonium
15 acetate, methyltridecylammonium propionate,
methyltridecylammonium butyrate, methyltridecylammonium
isobutyrate, methyltridecylammonium valerate,
methyltridecylammonium pivalate, methyltridecylammonium
hexanoate, methyltridecylammonium octanoate,
20 methyltridecylammonium decanoate, methyltridecylammonium
lactate, tetraethylammonium formate, tetraethylammonium
acetate, tetraethylammonium propionate, tetraethylammonium
butyrate, tetraethylammonium isobutyrate, tetraethylammonium
valerate, tetraethylammonium pivalate, tetraethylammonium
25 hexanoate, tetraethylammonium octanoate, tetraethylammonium

decanoate, tetraethylammonium lactate, tetrapropylammonium formate, tetrapropylammonium acetate, tetrapropylammonium propionate, tetrapropylammonium butyrate, tetrapropylammonium isobutyrate, tetrapropylammonium valerate,

5 tetrapropylammonium pivalate, tetrapropylammonium hexanoate, tetrapropylammonium octanoate, tetrapropylammonium decanoate, tetrapropylammonium lactate, tetrabutylammonium formate, tetrabutylammonium acetate, tetrabutylammonium propionate, tetrabutylammonium butyrate, tetrabutylammonium isobutyrate,

10 tetrabutylammonium valerate, tetrabutylammonium pivalate, tetrabutylammonium hexanoate, tetrabutylammonium octanoate, tetrabutylammonium decanoate, tetrabutylammonium lactate, tetramethylammonium methyl sulfate, tetraethylammonium, methyl sulfate, tetrapropylammonium methyl sulfate,

15 tetrabutylammonium methyl sulfate, tetramethylammonium methanesulfate, tetraethylammonium methanesulfate, tetrapropylammonium methasulfate, tetrabutylammonium methasulfate, tetramethylammonium ethanesulfate, tetrabutylammonium ethanesulfate, tetraethylammonium ethanesulfate, tetrapropylammonium

20 ethanesulfate, tetrabutylammonium ethanesulfate, tetramethylammonium benzenesulfate, tetraethylammonium benzenesulfate, tetrapropylammonium benzenesulfate, tetrabutylammonium benzenesulfate, tetramethylammonium toluenesulfate, tetraethylammonium toluenesulfate,

25 tetrapropylammonium toluenesulfate and tetrabutylammonium

toluenesulfate.

Moreover, the substance shown by general formula (2) may be quaternary ammonium sulfate, quaternary ammonium hydrogensulfate, quaternary ammonium sulfite, quaternary ammonium bisulfite, quaternary ammonium carbonate, quaternary ammonium hydrogencarbonate, quaternary ammonium oxalate, quaternary ammonium hydrogenoxalate, quaternary ammonium nitrate or quaternary ammonium nitrite which comprises a substituent represented by the abovementioned R¹.

The quaternary ammonium salt represented by general formula (2) may include bistetramethylammonium sulfate, bistetraethylammonium sulfate, bistetrapropylammonium sulfate, bistetrabutylammonium sulfate, tetramethylammonium hydrogensulfate, tetraethylammonium hydrogensulfate, tetrapropylammonium hydrogensulfate, tetrabutylammonium hydrogensulfate, bistetramethylammonium sulfite, bistetraethylammonium sulfite, bistetrapropylammonium sulfite, bistetrabutylammonium sulfite, tetramethylammonium hydrogensulfite, tetraethylammonium hydrogensulfite, tetrapropylammonium hydrogensulfite, tetrabutylammonium hydrogensulfite, bistetramethylammonium carbonate, bistetraethylammonium carbonate, bistetrapropylammonium carbonate, bistetrabutylammonium carbonate, tetramethylammonium hydrogencarbonate, tetraethylammonium hydrogencarbonate, tetrapropylammonium hydrogencarbonate,

tetrabutylammonium hydrogencarbonate, bistetramethylammonium oxalate, bistetraethylammonium oxalate, bistetrapropylammonium oxalate, bistetrabutylammonium oxalate, tetramethylammonium hydrogenoxalate, tetraethylammonium 5 hydrogenoxalate, tetrapropylammonium hydrogenoxalate, tetrabutylammonium hydrogenoxalate, tetramethylammonium nitrate, tetraethylammonium nitrate, tetrapropylammonium nitrate and tetrabutylammonium nitrate.

The quaternary ammonium salt in this case may be added 10 in an amount of 0.001 to 10 parts by weight, more preferably 0.005 to 5 parts by weight, still more preferably 0.01 to 1 part by weight per 1 part by weight of the siloxane polymer. After the addition, stirring between 10 minutes and 10 hours at 0 to 100°C under 10 Pa to 5 MPa can provide a target 15 siloxane polymer whose frame structure is changed. In addition, the solvent for having dissolved the siloxane polymer can be replaced by the solvent used for the coating liquid at this stage.

The solvent to be used for the coating liquid may 20 include aliphatic hydrocarbon such as n-pentane, isopentane, n-hexane, isohexane, n-heptane, 2,2,4-trimethylpentane, n-octane, isoctane, cyclohexane and methylcyclohexane; aromatic hydrocarbon such as benzene, toluene, xylene, ethylbenzene, trimethylbenzene, methylethylbenzene, n- 25 propylbenzene, isopropylbenzene, diethylbenzene,

isobutylbenzene, triethylbenzene, diisopropylbenzene and n-
amyl naphthalene; ketone such as acetone, methyl ethyl ketone,
methyl n-propyl ketone, methyl n-butyl ketone, methyl
isobutyl ketone, cyclohexanone, 2-hexanone,
5 methylcyclohexanone, 2,4-pentanedione, acetonyl acetone,
diacetone alcohol, acetophenone and fenthion; ether such as
ethyl ether, isopropyl ether, n-butyl ether, n-hexyl ether,
2-ethylhexyl ether, dioxolane, 4-methyldioxolane, dioxane,
dimethyl dioxane, ethylene glycol mono-n-butyl ether,
10 ethylene glycol mono-n-hexyl ether, ethylene glycol
monophenyl ether, ethylene glycol mono-2-ethyl butyl ether,
ethylene glycol dibutyl ether, diethylene glycol monomethyl
ether, diethylene glycol dimethyl ether, diethylene glycol
monoethyl ether, diethylene glycol dimethyl ether, diethylene
15 glycol monoethyl ether, diethylene glycol diethyl ether,
diethylene glycol monopropyl ether, diethylene glycol
dipropyl ether, diethylene glycol monobutyl ether, diethylene
glycol dibutyl ether, tetrahydrofuran, 2-
methyltetrahydrofuran, propylene glycol monomethyl ether,
20 propylene glycol dimethyl ether, propylene glycol monoethyl
ether, propylene glycol diethyl ether, propylene glycol
monopropyl ether, propylene glycol dipropyl ether, propylene
glycol monobutyl ether, dipropylene glycol dimethyl ether,
dipropylene glycol diethyl ether, dipropylene glycol dipropyl
ether and dipropylene glycol dibutyl ether; ester such as
25

diethyl carbonate, ethyl acetate, γ -butyrolactone, γ -valerolactone, n-propyl acetate, isopropyl acetate, n-butyl acetate, isobutyl acetate, sec-butyl acetate, n-pentyl acetate, 3-methoxybutyl acetate, methylpentyl acetate, 2-5 ethylbutyl acetate, 2-ethylhexyl acetate, benzyl acetate, cyclohexyl acetate, methylcyclohexyl acetate, n-nonyl acetate, methyl acetoacetate, ethyl acetoacetate, ethylene glycol monomethyl ether acetate, ethylene glycol monoethyl ether acetate, diethylene glycol monomethyl ether acetate, 10 diethylene glycol monoethyl ether acetate, diethylene glycol mono-n-butyl ether acetate, propylene glycol monomethyl ether acetate, propylene glycol monoethyl ether acetate, dipropylene glycol monomethyl ether acetate, dipropylene glycol monoethyl ether acetate, dipropylene glycol mono-n-15 butyl ether acetate, glycol diacetate, methoxy triglycol acetate, ethyl propionate, n-butyl propionate, isoamyl propionate, diethyl oxalate, di-n-butyl oxalate, methyl lactate, ethyl lactate, n-butyl lactate, n-amyl lactate, diethyl malonate, dimethyl phthalate and diethyl phthalate; 20 nitrogen-containing solvent such as N-methylformamide, N,N-dimethylformamide, acetamide, N-methylacetamide, N,N-dimethylacetamide, N-methylpropionamide and N-methylpyrrolidone; and sulfur-containing solvent such as dimethyl sulfide, diethyl sulfide, thiophene, 25 tetrahydrothiophene, dimethylsulfoxide, sulfolane and 1,3-

propanesultone. It may be used singly or in an admixture of two or more.

Film having a desirable thickness can be obtained preferably by spin-coating with controlled concentration of 5 the solute at an appropriate speed of rotation. The actual thickness of the film may be, but not limited to, typically about 0.2 to 1 μm . However, membranes of different thicknesses can be made. For example, film having a large thickness can be made by repeating the coating. The solvent 10 to be used for diluting the coating liquid may include same solvent as the replacement solvent for the coating liquid. It can be used singly or in an admixture of two or more.

Although the level of the dilution may depend on the 15 viscosity or the desired film thickness, the typical content of the solvent or solvents may be 50 to 99% by weight, more preferably 75 to 95% by weight.

The obtained film is heated. This is a heating process (usually called a 'prebake' in a semiconductor process) for 20 drying the coated film. The film may be heated preferably at 50 to 150°C for several minutes so as to remove the solvent. After the heating step for drying, another heating step for curing the coated film is provided. In the heating step for curing, the coating film may be heated preferably at 150 to 25 500°C, more preferably 180 to 450°C. The heating time may be

preferably 1 to 300 minutes, more preferably 1 to 100 minutes. The heating step for drying and the heating step for curing may be carried out in one heating process or in two heating processes. The obtained film has high mechanical strength in 5 the entire film and may have typically 0.7 to 5 GPa of hardness and 5 to 40 GPa of modulus of elasticity according to the measurement by the nanoindentation. On the other hand, according to the measurement of usual porous material having pores produced thermally by removing the thermally 10 decomposable polymers which have been contained in the silicone resin, the hardness is 0.05 to 2 Gpa and the modulus of elasticity is about 1.0 to 4.0 GPa. Thus, the mechanical strength of the film of the invention is extremely high.

The porous film of the present invention is 15 particularly preferable as the interlevel insulator film of the interconnections in a semiconductor integrated circuit. The semiconductor device is required to reduce interconnection capacitance in order to prevent interconnection delay when highly integrated. Various means 20 have been developed to achieve this, and one of them is to reduce the relative permittivity (dielectric constant) of the interlevel insulator film disposed between metal interconnections. When an interlevel insulator film is prepared by using the composition for forming a porous film 25 of the present invention, the semiconductor device can be

downsized and faster and consume less power.

However, conventionally, there is a problem that when a porous film is prepared by introducing pores in the film so as to lower the dielectric constant, the mechanical strength 5 of the film decreases as the density of the material composing the film decreases. The decrease in mechanical strength not only affects the strength of the semiconductor device itself but also causes exfoliation due to insufficient strength in a chemical mechanical polishing process, which is 10 generally used in the fabrication process. Particularly, when used as the interlevel insulator film of a semiconductor, the porous film of the invention having high mechanical strength despite the porosity prevents such exfoliation. Consequently, a highly reliable semiconductor having high speed performance 15 in a small size can be provided.

The embodiments of the semiconductor device of the present invention will be described below. Fig. 1 shows a schematic cross-sectional view of an example of the semiconductor device of the present invention.

20 In Fig. 1, the substrate 1 is an Si semiconductor substrate such as an Si substrate or an SOI (Si-on-insulator) substrate; however, it can be a compound semiconductor substrate such as SiGe or GaAs. The interlevel insulator films include the interlevel insulator film 2 of the contact 25 layer; the interlevel insulator films 3, 5, 7, 9, 11, 13, 15,

and 17 of the interconnection layers; and the interlevel insulator films 4, 6, 8, 10, 12, 14, and 16 of the via layers. The interconnection layers corresponding to the lowermost interlevel insulator film 3 through the uppermost insulator film 17 are abbreviated as M1, M2, M3, M4, M5, M6, M7, and M8, respectively. The via layers corresponding to the lowermost interlevel insulator film 4 through the uppermost insulator film 16 are abbreviated as V1, V2, V3, V4, V5, V6, and V7, respectively. Although some of the metal interconnections are referred to with the numbers 18 and 21 to 24, the other regions with the same pattern not labeled with numbers indicate metal interconnections. The via plug 19 is made from a metal. In the case of copper interconnection, copper is generally used. The regions having the same pattern as the via plug 19 represent via plugs although they are not labeled with numbers in the drawing. The contact plug 20 is connected to the gate of the transistor (not illustrated) formed on the top surface of the substrate 1 or to the substrate. Thus, the interconnection layers and the via layers are alternately stacked, and multilayer interconnections generally indicate M1 and regions higher than M1. In general, M1 to M3 are called local interconnections, M4 and M5 are called intermediate interconnections or semi-global interconnections, and M6 to M8 are called global interconnections.

In the semiconductor device of the present invention, the porous film of the present invention is used as one or more of the interlevel insulator films 3, 5, 7, 9, 11, 13, 15, and 17 of the interconnection layers or the insulator films 4, 5 6, 8, 10, 12, 14, and 16 of the via layers.

For example, when the porous film of the present invention is used for the interlevel insulator film 3 of the interconnection layer (M1), the interconnection capacitance between the metal interconnection 21 and the metal 10 interconnection 22 can be greatly reduced. When the porous film of the present invention is used for the interlevel insulator film 4 of the via layer (V1), the interconnection capacitance between the metal interconnection 23 and the metal interconnection 24 can be greatly reduced. Using the 15 porous film with a low relative permittivity of the present invention as an interconnection layer can greatly reduce the metal interconnection capacitance in the same layer. On the other hand, using the porous film with a low relative 20 permittivity of the present invention as a via layer can greatly reduce the capacitance between the upper and lower metal interconnection layers.

Therefore, using the porous film of the present invention for all of the interconnection layers and the via layers can greatly reduce the parasitic capacitance of the 25 interconnections. The porous film of the present invention

enables a semiconductor device to have higher mechanical strength by its high mechanical strength, thereby greatly improving the yield of the fabrication and the reliability of the semiconductor device.

5 The invention will be described specifically through the following examples, but is not limited to them.

Synthetic Example 1 (synthesis of siloxane polymer with acidic catalyst)

An aqueous 0.05 wt% nitric acid solution of 60 g was
10 added to the solution mixture of tetraethoxysilane (80g), methyltrimethoxysilane (50g) and ethanol (460g), and stirred for four hours at 25°C. The ethanolic solution of the silicone polymer was obtained.

15 Synthetic Example 2. (synthesis of siloxane polymer with basic catalyst)

A solution of tetraethoxysilane (60g) and methyltrimethoxysilane (30g) was added to the mixture of the solution of an aqueous 40wt% methylamine solution (10g),
20 ultra pure water (640g) and ethanol (1200g), and stirred for four hours at 75°C. The ethanolic solution of the silicone polymer was obtained.

Synthesis Example 3 (synthesis of siloxane polymer with basic catalyst)

A solution of tetramethoxysilane (40g) and methyltrimethoxysilane (30g) was added to the mixture of an aqueous 28wt% ammonia solution (10g), ultra pure water (640g) and ethanol (1200g), and stirred for four hours at 75°C. The 5 ethanolic solution of silicone polymer was obtained.

Example 1

An aqueous 5 wt% tetramethylammonium caproate solution (90g) and propylene glycol monopropyl ether (700g) were added 10 at 25°C to the siloxane polymer solution (650 g) obtained in Synthetic Example 1. After stirring for one hour, the reaction mixture was concentrated under reduced pressure at 40°C so that the coating liquid (700g) was obtained. Then, it was applied on an eight-inch wafer with a spin coater for 15 one minute at 1,500 rpm. The film thickness, when heated for two minutes at 120°C using a hot plate, was 8,000A. After it was further heated at 250°C for three minutes, it was heated at 450°C for one hour in a nitrogen atmosphere in a clean oven. The thickness was 7,200A. The dielectric constant of 20 the obtained film was 2.2 and the modulus of elasticity thereof was 6.5 Gpa. The coating liquid obtained was left stand at 23°C for 30 days. Afterwards, the film was obtained in the same manner as the above. Then, the values of the properties were obtained.

Method for measuring properties

<dielectric constant>

It was measured by CV technique using an automatic mercury probe with an automatic mercury CV measurement device 5 495-CV system (made by SSM Japan KK).

<modulus of elasticity>

It was measured with a Nano indenter (made by the Nano Instruments Innovation Center).

10 Example 2

An aqueous 5wt% trimethyloctylammonium acetate solution (16g) and propylene glycol monopropyl ether (300g) were added at 25°C to the solution (1kg) obtained in Synthetic Example 2. After it was stirred for one hour, the reaction mixture was 15 concentrated under reduced pressure at 40°C and the coating liquid (300g) was obtained. The film was then formed in the same manner as that in Example 1. The dielectric constant of the coated film formed in the same manner was 2.1, and the modulus of elasticity was 5.5 GPa. The properties were 20 measured after 30 days in same manner as in Example 1, the identical values were obtained.

Example 3

An aqueous 5wt% tetrapropylammonium acetate solution 25 (16g) and propylene glycol monopropyl ether (200g) were added

at 25°C to the solution (1kg) obtained in Synthetic Example 3. After it was stirred for one hour, the reaction mixture was concentrated under reduced pressure at 40°C and the coating liquid (200g) was obtained. Then the film was formed in the 5 same manner in Example 1. The dielectric constant of the coated film formed in the same manner was 2.2 and the modulus of elasticity was 6.3 GPa. The properties were measured after 30 days in the same manner as in Example 1, the identical values were obtained.

10

Example 4

An aqueous 10 wt% bistetramethylammonium carbonate solution (45g) and propylene glycol monopropyl ether (700g) were added at 25°C to the siloxane polymer solution (650g) 15 obtained in Synthetic Example 1. After stirring for one hour at 40°C, the reaction mixture was concentrated under reduced pressure at 40°C and the coating liquid (700g) was obtained. The film was then formed in the same manner as in Example 1. The dielectric constant of the coated film formed in the same 20 manner was 2.4 and the modulus of elasticity was 8.7GPa. The properties were measured after 30 days in the same manner as in Example 1, the exactly same values were obtained.

Example 5

25 An aqueous 5wt% tetramethylammonium p-toluenesulfonate

solution (90g) and propylene glycol monopropyl ether (700g) were added at 25°C to the siloxane polymer solution (650g) obtained in Synthetic Example 1. After it was stirred for one hour, the reaction mixture was concentrated under reduced pressure at 40°C and the coating liquid (700g) was obtained.

5 The film was then formed in the same manner as in Example 1. The dielectric constant of the coated film formed in the same manner was 2.5 and the modulus of elasticity was 7.0 GPa. The properties were measured after 30 days in the same manner

10 in Example 1 and exactly same values were obtained.

Example 6

An aqueous 5 wt% tetrabutylammonium acetate solution (16g) and propylene glycol monopropyl ether (300g) were added 15 25°C to the solution (1kg) obtained in Synthetic Example 2. After it was stirred at 40°C for one hour, the reaction mixture was concentrated under reduced pressure at 40°C and the coating liquid (300g) was obtained. The film was then formed in the same manner in Example 1. The dielectric constant of the coated film formed in the same manner was 2.1 20 and the modulus of elasticity was 5.1 GPa. The properties were measured after 30 days in the same manner in Example 1, the exactly same values were obtained.

Example 7

An aqueous 10wt% tetramethylammonium sulfate solution (13g) and propylene glycol monopropyl ether (300g) were added at 25°C to the solution (1kg) obtained in Synthetic Example 2. After it had been stirred for one hour, the reaction mixture 5 was concentrated under reduced pressure at 40°C and the coating liquid (300g) was obtained. The film was then formed in the same manner as in Example 1. The dielectric constant of the coated film formed in the same manner was 2.2 and the modulus of elasticity was 5.8GPa. The properties were 10 measured after 30 days in the same manner as in Example 1 and the exactly same values were obtained.

Example 8

An aqueous 5wt% tributylmethylammonium formate solution 15 (25g) and propylene glycol monopropyl ether (200g) were added at 25°C to the solution (1kg) obtained in Synthetic Example 3. After it was stirred for one hour, the reaction mixture was concentrated under reduced pressure at 40°C and the coating liquid (400g) was obtained. The film was then formed in the 20 same manner as in Example 1. The dielectric constant of the coated film formed in the same manner was 2.0 and the modulus of elasticity was 4.3 GPa. The properties were measured after 30 days in the same manner as in Example 1, the exact same values were obtained.

Example 9

An aqueous 10wt% tetraethylammonium lactate solution and propylene glycol monopropyl ether (250g) were added at 25°C to the solution (1kg) obtained in Synthetic Example 3. After 5 it was stirred for one hour, the reaction mixture was concentrated under reduced pressure at 40°C and the coating liquid (250g) was obtained. The film was then formed in a similar way to that in Example 1. The dielectric constant of the coated film formed in the same manner was 2.4 and the 10 modulus of elasticity was 5.3 GPa. The properties were measured after 30 days in the same manner in Example 1 and the exactly same values were obtained.

Comparative Example 1

15 Experiment was carried out in the same manner as in In Example 1, except that the aqueous 5wt% tetramethylammonium hydroxide solution (90 g) was added instead of the aqueous 5wt% tetramethylammonium caproate solution. Then the coated film having dielectric constant of 2.4 and modulus of 20 elasticity of 3.3 GPa was obtained. When the coating liquid was left stand at 23°C, it gelated in three days so that the coating application became impossible.

Comparative Example 2

25 The experiment was carried in the same manner in Example

2 except that an aqueous 5wt% trimethyloctylammonium hydroxide solution (16g) was added instead of the aqueous 5wt% trimethyloctylammonium acetate solution in Example 2. Then, coated film having dielectric constant of 2.3 and 5 modulus of elasticity of 1.9 GPa was obtained. When the coating liquid was left stand at 23°C, it gelated in three days so that the coating application became impossible.

Comparative Example 3

10 The experiment was carried in the same manner in Example 3 except an aqueous 5wt% tetrapropylammonium bromide (16g) was added instead of the aqueous 5wt% tetrapropylammonium acetate solution. Then, coated film having dielectric constant of 2.4 and modulus of elasticity of 1.8GPa was 15 obtained. The 50ppm bromine was detected in the film so that corrosion of the wiring material was possible. Therefore, it became evident that it was useless for actual semiconductor device manufacturing.

20 The results are summarized in Table 1.

Table 1

	siloxane polymer	quaternary ammonium salt	dielectric constant	modulus (GPa)	remark
Example1	Syn. Ex. 1	$(\text{CH}_3)_4\text{N}^+(\text{C}_5\text{H}_{11}\text{COO})^-$	2.2	6.5	no property change after 30days
Example2	Syn. Ex. 2	$[(\text{CH}_3)_3(\text{C}_8\text{H}_{17})\text{N}]^+(\text{CH}_3\text{COO})^-$	2.1	5.5	no property change after 30days
Example3	Syn. Ex. 3	$(\text{C}_3\text{H}_7)_4\text{N}^+(\text{CH}_3\text{COO})^-$	2.2	6.3	no property change after 30days
Example4	Syn. Ex. 1	$[(\text{CH}_3)_4\text{N}^+]_2(\text{CO}_3)^{2-}$	2.4	8.7	no property change after 30days
Example5	Syn. Ex. 1	$(\text{CH}_3)_4\text{N}^+(\text{p}-\text{CH}_3-\text{ph}-\text{SO}_3)^-$	2.5	7.0	no property change after 30days
Example6	Syn. Ex. 2	$(\text{C}_4\text{H}_9)_4\text{N}^+(\text{CH}_3\text{COO})^-$	2.1	5.1	no property change after 30days
Example7	Syn. Ex. 2	$[(\text{CH}_3)_4\text{N}^+]_2(\text{SO}_4)^{2-}$	2.2	5.8	no property change after 30days
Example8	Syn. Ex. 3	$[(\text{C}_4\text{H}_9)_3(\text{CH}_3)\text{N}]^+(\text{HCOO})^-$	2.0	4.3	no property change after 30days
Example9	Syn. Ex. 3	$(\text{C}_2\text{H}_5)_4\text{N}^+[\text{CH}_3\text{CH}(\text{OH})\text{COO}]^-$	2.4	5.3	no property change after 30days
Comp. Ex. 1	Syn. Ex. 1	$(\text{CH}_3)_4\text{N}^+(\text{OH})^-$	2.4	3.3	gelation after 3 days
Comp. Ex. 2	Syn. Ex. 2	$[(\text{CH}_3)_3(\text{C}_8\text{H}_{17})\text{N}]^+(\text{OH})^-$	2.3	1.9	gelation after 5 days
Comp. Ex. 3	Syn. Ex. 3	$(\text{C}_3\text{H}_7)_4\text{N}^+(\text{Br})^-$	2.4	1.8	bromine porion 50ppm